

K800844 MEDEBAR XLMay 30, 1980
45 days to decisionK800844 · Product code: **KTA** · Radiology
Source: <https://www.510kdatabase.net/k800844/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Medium, Contrast, Radiologic (KTA)
Date received	Apr 15, 1980
Decision date	May 30, 1980
Days to decision	45 days
Third-party review	No

APPLICANT

Company	Medefield Pty. , Ltd.
Location	Mchenry, IL, US
510(k) history	6 submissions · 6 cleared · 1980-1980

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k800844/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 22, 2026